APPENDIX A Text of the Relevant Claims in the '793 and '782 Patents

Claims 1 and 4 of the '793 Patent Claim 1 of the '782 Patent 1. A method of treating pulmonary hypertension comprising: providing an inhalation device for treating pulmonary hypertension in a human suffering from pulmonary hypertension comprising a powder formulation of treprostinil or a pharmaceutically acceptable salt thereof and a dry powder inhaler configured to administer single event dose of the powder formulation comprising treprostinil or a pharmaceutically acceptable salt thereof, wherein the single event dose 1. A method of treating pulmonary hypertension compriscomprises at least 15 micrograms to 90 micrograms of ing administering by inhalation to a human suffering from treprostinil or a pharmaceutically acceptable salt thereof pulmonary hypertension a therapeutically effective single delivered in 1 to 3 breaths, wherein the dry powder inhaler event dose of a formulation comprising treprostinil or a is configured to administer the entire single event dose in pharmaceutically acceptable salt thereof with an inhalation less than 5 minutes with at least 5 micrograms of treprostinil device, wherein the therapeutically effective single event or a pharmaceutically acceptable salt thereof being inhaled dose comprises from 15 micrograms to 90 micrograms of per breath through coordinated actuation of the dry powder treprostinil or a pharmaceutically acceptable salt thereof inhaler with each breath, and administering to a human delivered in 1 to 3 breaths. suffering from pulmonary hypertension with the dry powder inhaler the single event dose comprising at least 15 micro-4. The method of claim 1, wherein the inhalation device grams to 90 micrograms of treprostinil or a pharmaceutically is a dry powder inhaler. acceptable salt thereof in 1 to 3 breaths, wherein the human administers the entire single event dose with the dry powder inhaler in less than 5 minutes by inhaling at least 5 micrograms of treprostinil or a pharmaceutically acceptable salt thereof per breath by coordinating one actuation of the dry powder inhaler for each separate breath, and wherein administration of an additional single event dose in the same manner occurs at least 3 hours later.